

Marcia L E Macdonald

List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

2,116
citations

1040056

9
h-index

1281871

11
g-index

12
all docs

12
docs citations

12
times ranked

2487
citing authors

#	ARTICLE	IF	CITATIONS
1	The metabolic phenotype of SCD1-deficient mice is independent of melanin-concentrating hormone. <i>Peptides</i> , 2010, 31, 123-129.	2.4	2
2	Despite Antiatherogenic Metabolic Characteristics, SCD1-Deficient Mice Have Increased Inflammation and Atherosclerosis. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2009, 29, 341-347.	2.4	95
3	Absence of stearyl-CoA desaturase-1 does not promote DSS-induced acute colitis. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2009, 1791, 1166-1172.	2.4	12
4	Absence of stearyl-CoA desaturase-1 ameliorates features of the metabolic syndrome in LDLR-deficient mice. <i>Journal of Lipid Research</i> , 2008, 49, 217-229.	4.2	59
5	Loss of function mutations in the Na ^v 1.7 gene underlie congenital indifference to pain in multiple human populations. <i>Clinical Genetics</i> , 2007, 71, 311-319.	2.0	404
6	Ironing out neurodegeneration: mutations in a phospholipase A2 cause neurodegenerative diseases with iron accumulation. <i>Clinical Genetics</i> , 2006, 70, 306-307.	2.0	4
7	FASA-57 cDNA shares no homology with coding sequence of HD gene. <i>Journal of Reproductive Immunology</i> , 2006, 69, 9-10.	1.9	0
8	Genetic variants of frizzled-4 gene in familial exudative vitreoretinopathy and advanced retinopathy of prematurity. <i>Clinical Genetics</i> , 2005, 67, 363-366.	2.0	49
9	Mutations in HFE2 cause iron overload in chromosome 1q-linked juvenile hemochromatosis. <i>Nature Genetics</i> , 2004, 36, 77-82.	21.4	900
10	Identification of a Novel Gene (HSN2) Causing Hereditary Sensory and Autonomic Neuropathy Type II through the Study of Canadian Genetic Isolates. <i>American Journal of Human Genetics</i> , 2004, 74, 1064-1073.	6.2	133
11	Identification and functional analysis of a naturally occurring E89K mutation in the ABCA1 gene of the WHAM chicken. <i>Journal of Lipid Research</i> , 2002, 43, 1610-1617.	4.2	49
12	Mutant frizzled-4 disrupts retinal angiogenesis in familial exudative vitreoretinopathy. <i>Nature Genetics</i> , 2002, 32, 326-330.	21.4	409